Title 21—Food and Drugs

(This book contains parts 600 to 799)

CHAPTER I—Food and Drug Administration, Department of Health and Human Services (Continued)	60	0
CROSS REFERENCES: Food Safety and Inspection Service, Department of Agriculture: CFR Chapter III. Federal Trade Commission: See Commercial Practices, 16 CFR Chapter I	See	9

Part

U.S. Customs Service, Department of the Treasury: See Customs Duties, 19 CFR Chapter I.

Internal Revenue Service, Department of the Treasury: See Internal Revenue, 26 CFR Chapter I

Bureau of Alcohol, Tobacco, and Firearms, Department of the Treasury: See Alcohol, Tobacco Products and Firearms, 27 CFR Chapter I.

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH AND HUMAN SERVICES (CONTINUED)

(Parts 600-799)

EDITORIAL NOTE: For nomenclature changes to chapter I see 59 FR 14366, Mar. 28, 1994, and 66 FR 56035, Nov. 6, 2001.

SUBCHAPTER F—BIOLOGICS

Part		Page
600	Biological products: general	5
601	Licensing	19
606	Current good manufacturing practice for blood and blood components	42
607	Establishment registration and product listing for manufacturers of human blood and blood products	54
610	General biological products standards	60
630	General requirements for blood, blood components, and blood derivatives	84
640	Additional standards for human blood and blood products	85
660	Additional standards for diagnostic substances for laboratory tests	109
680	Additional standards for miscellaneous products	122
	SUBCHAPTER G—COSMETICS	
700	General	126
701	Cosmetic labeling	132
710	Voluntary registration of cosmetic product establishments	145
720	Voluntary filing of cosmetic product ingredient composition statements	146
740	Cosmetic product warning statements	150
741–799	[Reserved]	